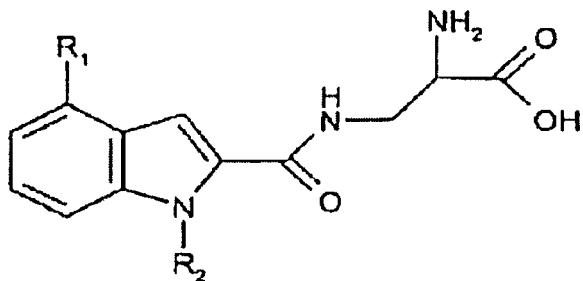


## CLAIMS

1. (Cancelled)
2. (Currently amended) A compound of formula I



wherein

R<sub>1</sub> is phenyl or naphthyl, wherein phenyl is substituted by one or two of halogen, C<sub>1-6</sub>-alkyl, C<sub>1-6</sub>-alkoxy or phenyl-C<sub>1-6</sub>-alkyl; and  
R<sub>2</sub> is hydrogen or C<sub>1-6</sub>-alkyl;  
in free, hydrate or salt form.

3. (Currently amended) A compound according to Claim 4 which that is selected from 3-(4-(2-ethylphenyl)-2-carboxamido-indole)-alanine, 3-(4-(2-benzyl-phenyl)-2-carboxamido-indole)-alanine, 3-(4-(naphthalen-2-yl)-2-carboxamido-indole)-alanine, 3-(4-(naphthalen-1-yl)-2-carboxamido-indole)-alanine, 3-(4-(2-butoxy-phenyl)-2-carboxamido-indole)-alanine, 3-(4-(2-propyl-phenyl)-2-carboxamido-indole)-alanine, 3-(4-(2-isopropyl-phenyl)-2-carboxamido-indole)-alanine, and 3-(4-(2,4-dichloro-phenyl)-2-carboxamido-indole)-alanine, or a and pharmaceutically-acceptable salts therefor thereof.

4. (Currently amended) A compound according to Claim 3 which that is 3-(4-(2-ethylphenyl)-2-carboxamido-indole)-D-alanine, in free form or in a pharmaceutically-acceptable salt form.

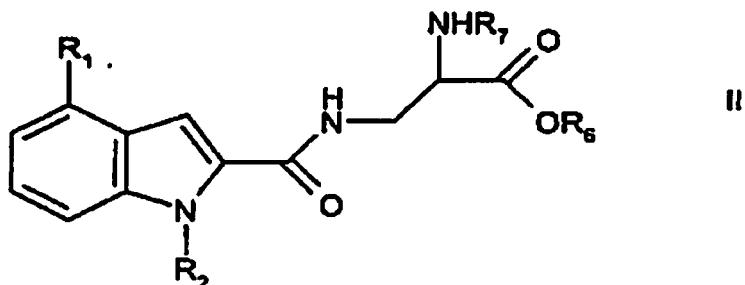
5. (Canceled)

6. (Currently amended) A pharmaceutical composition comprising a compound as defined according to Claim 4\_2, in free form or in pharmaceutically-acceptable salt form, in association with a pharmaceutically-acceptable diluent or carrier therefor.

7. (Canceled)

8. (Currently amended) A pharmaceutical combination comprising a compound according to Claim 4\_2 in free form or in a pharmaceutically-acceptable salt form and a further agent selected from immunosuppressant, immunomodulatory, anti-inflammatory and chemotherapeutic drug agents.

9. (Withdrawn by the Examiner) A process for the production of the compound according to claim 2, which process comprises deprotecting a compound of



wherein R<sub>1</sub> and R<sub>2</sub> are as defined in claim 2,

R<sub>6</sub> is C<sub>1-6</sub>alkyl or benzyl,

R<sub>7</sub> is an amino protecting group,

and optionally converting the compound of formula I obtained in free form to a salt form or vice versa.

10. (Withdrawn by the Examiner) A method for treating or preventing disorders or diseases mediated by lymphocytes, acute or chronic transplant rejection, T-

cell mediated inflammatory or autoimmune diseases, diabetes, allergic diseases, myocarditis, hepatitis, ischemia/reperfusion injury, renal failure, hemorrhage shock, traumatic shock, cancer or infectious diseases, in a subject in need of such treatment, which method comprises administering to said subject an effective amount of a compound according to claim 1, or a pharmaceutically acceptable salt thereof.